What is claimed is:

1. A compound for modulating kinase activity, particularly Tie-2, of Formula I,

$$X = \begin{bmatrix} R^1 & R^3 & & & \\ & R^2 & & & \\ & & & \\ & & & & \\ & & &$$

5 or a pharmaceutically acceptable salt, hydrate, or prodrug thereof, wherein,

X is selected from -H, -OR⁶, -S(O)₀₋₂R⁶, -N(R⁶)R⁷, -O-N(R⁶)R⁷, -N(R⁶)OR⁶, -N(R⁶)N(R⁶)R⁷, absent, oxo, thiono, and imino, with the proviso that when X is oxo, thiono, or imino, there is only one R^1 ;

R¹ and R², at each occurance, are each independently selected from -H, halogen, -CN, -NH₂, -NO₂, -OR⁶, -N(R⁶)R⁷, -S(O)₀₋₂R⁷, -SO₂N(R⁶)R⁷, -CO₂R⁶, -C(O)N(R⁶)R⁷, -N(R⁶)SO₂R⁷, -N(R⁶)CO₂R⁷, -N(R⁶)CO₂R⁷, -C(O)R⁶, optionally substituted lower alkyl, optionally substituted aryl, optionally substituted lower arylalkyl, optionally substituted heterocyclyl, absent, and optionally substituted lower heterocyclylalkyl;

optionally two of R² together are oxo;

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optionally, at least one pair of substituents, selected from two of R¹, two of R², and one each of R¹ and R², together with the corresponding carbon or carbons to which they are attached, form a first ring comprising between three and seven annular atoms, said first ring optionally substituted with between zero and four additional of R¹, each independently selected as defined above and optionally, when paired, together with the corresponding atom or atoms of the first ring to which they are attached, form a second ring comprising between three and seven annular atoms, said second ring optionally substituted with between zero and three of R¹;

R³ is selected from -H, optionally substituted lower alkyl, optionally substituted lower arylalkyl, optionally substituted aryl, optionally substituted heterocyclyl, and optionally substituted alkoxy;

optionally R³ and one of R², together with the atoms to which each is attached, form a third ring comprising between three and seven annular atoms, said third ring optionally

substituted with between zero and four additional of R¹, each independently selected as defined above and optionally, when paired, together with the corresponding atom or atoms of the third ring to which they are attached, form a fourth ring comprising between three and seven annular atoms, said fourth ring optionally substituted with between zero and three of R¹;

optionally R³ and one of R¹, together with the atoms to which they are attached and the carbon to which R² is attached, form a fifth ring comprising between three and seven annular atoms atoms, said fifth ring optionally substituted with between zero and four additional of R¹, each independently selected as defined above and optionally, when paired, together with the corresponding atom or atoms of the fifth ring to which they are attached, form a sixth ring comprising between three and seven annular atoms, said sixth ring optionally substituted with between zero and three of R¹;

m is zero to four;

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each of R^4 is independently selected from -H, halogen, -CN, -NH₂, -NO₂, -OR⁶, -N(R⁶)R⁷, -S(O)₀₋₂R⁷, -SO₂N(R⁶)R⁷, -CO₂R⁶, -C(O)N(R⁶)R⁷, -N(R⁶)SO₂R⁷, -N(R⁶)C(O)R⁷, -N(R⁶)CO₂R⁷, -C(O)R⁶, optionally substituted lower alkyl, optionally substituted aryl, optionally substituted lower arylalkyl, optionally substituted heterocyclyl, and optionally substituted lower heterocyclylalkyl;

optionally two adjacent of R^4 , together with the two carbons to which they are attached, form a seventh ring fused with the aromatic ring system containing Z as in Formula I, said seventh ring comprising between five and seven atoms and substituted with zero to three additional of R^4 , provided said seventh ring together with the aromatic ring system containing Z as in Formula I does not constitute a 7-deazapurine;

each Y is independently either $=C(R^5)$ - or =N-, provided that there are no more than three of =N- in the aromatic ring bearing Y;

each Z is independently either $=C(R^4)$ - or =N-;

n is zero to five;

each R^5 is independently selected from -H, halogen, -CN, -NH₂, -NO₂, -OR⁶, -NR⁶R⁷, -S(O)₀₋₂R⁷, -SO₂NR⁶R⁷, -CO₂R⁶, -C(O)NR⁶R⁷, -N(R⁶)SO₂R⁷, -N(R⁶)C(O)R⁷, -N(R⁶)CO₂R⁷, -C(O)R⁶, optionally substituted lower alkyl, optionally substituted aryl, optionally substituted lower arylalkyl, optionally substituted heterocyclyl, and optionally substituted lower heterocyclylalkyl; and

optionally two adjacent of R⁵, together with the two carbons to which they are attached, form an eighth ring fused with the aromatic ring system containing Y as in Formula I, said eighth ring comprising between five and seven atoms and substituted with zero to three additional of R⁵;

5 R^6 is -H or R^7 ;

R⁷ is selected from optionally substituted lower alkyl, optionally substituted aryl, optionally substituted lower arylalkyl, optionally substituted heterocyclyl, and optionally substituted lower heterocyclylalkyl; and

R⁶ and R⁷, when taken together with a common nitrogen to which they are attached, form an optionally substituted five- to seven-membered heterocyclyl ring, said optionally substituted five- to seven-membered heterocyclyl ring optionally containing at least one additional heteroatom selected from N, O, S, and P.

2. The compound according to claim 1, of Formula II.

П

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- 3. The compound according to claim 2, wherein at least one of Z is =N-.
- 4. The compound according to claim 3, of Formula III.

$$R^{1}$$
 R^{1}
 R^{1}
 R^{3}
 R^{2}
 R^{2}
 R^{2}
 R^{4}
 R^{5})_r

5. The compound according to claim 4, wherein one each of R¹ and R², together with the corresponding carbons to which they are attached, form said first ring, said first ring

comprising a saturated ring, said saturated ring optionally substituted with between zero and four additional of R¹.

- 6. The compound according to claim 5, wherein said saturated ring is carbocyclic.
- 7. The compound according to claim 6, of Formula IV.

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$$(R^{1})_{1-4}$$
 $(R^{5})_{n}$
 $(R^{5})_{n}$
 $(R^{5})_{n}$

- 8. The compound according to claim 7, wherein two of R¹, together with the carbon or carbons to which they are attached, form said second ring.
- 9. The compound according to claim 8, wherein said second ring is a six-membered aryl, fused with said first ring, said second ring optionally substituted with between zero and three of R¹.
 - 10. The compound according to claim 9, of formula V.

$$(R^{1})_{0-3}$$
 $(R^{5})_{n}$
 $(R^{5})_{n}$

15 11. The compound according to claim 10, of formula VI.

$$(R^1)_{0-3}$$
 $(R^5)_r$
 $(R^4)_{0-3}$

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12. The compound according to claim 3, of formula VII,

$$(R^{1})_{0\cdot3} \xrightarrow{H} X \xrightarrow{R^{3}} N \xrightarrow{N} N \xrightarrow{N} (R^{5})_{0\cdot4}$$

$$VII$$

- 13. The compound according to claim 12, wherein at least one of R¹ is an optionally substituted aryl or an optionally substituted phenyl.
 - 14. The compound according to claim 12, wherein at least one of R^4 is an optionally substituted aryl or an optionally substituted phenyl.
 - 15. The compound according to claim 12, of formula VIII.

$$(R^{1})_{0-3}$$
 $(R^{5})_{0-4}$

10 **VIII**

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- 16. The compound according to claim 15, wherein two of R⁴, together with the aromatic annular atoms to which they are attached, form said seventh ring, said seventh ring comprising between zero and two nitrogens.
- 17. The compound according to claim 16, wherein said seventh ring is substituted with between zero and three additional of R⁴.
 - 18. The compound according to claim 1, selected from Table 3.

Table 3

#	Name	Structure
1	N-cyclohexyl-2-pyridin-4-ylquinazolin-4- amine	HN—Z

Table 3

#	Name	Structure
2	2-pyridin-4-yl-N-(2-pyrrolidin-1- ylethyl)quinazolin-4-amine	HN N
3	N-cyclopentyl-2-pyridin-4-ylquinazolin-4- amine	HN N
4	N-(cyclohexylmethyl)-2-pyridin-4- ylquinazolin-4-amine	
5	2-[(2-pyridin-4-ylquinazolin-4- yl)amino]ethanol	HN OH
6	3-[(2-pyridin-4-ylquinazolin-4- yl)amino]propan-1-ol	HN OH
7	N-[(4-fluorophenyl)methyl]-2-pyridin-4-ylquinazolin-4-amine	HN N F
8	N,N-dimethyl-N'-(2-pyridin-4-ylquinazolin-4-yl)ethane-1,2-diamine	HN N

Table 3

#	Name	Structure
9	N-(2,3-dihydro-1H-inden-1-yl)-2-pyridin- 4-ylquinazolin-4-amine	H Z Z Z
10	N-(2-morpholin-4-ylethyl)-2-pyridin-4- ylquinazolin-4-amine	HN N
11	4-[4-(2-pyridin-4-ylquinazolin-4- yl)piperazin-1-yl]phenol	HO————————————————————————————————————
12	2-pyridin-4-yl-N-[(2R)-1,2,3,4- tetrahydronaphthalen-2-yl]quinazolin-4- amine	
13	4-piperazin-1-yl-2-pyridin-4-ylquinazoline	
14	1,1-dimethylethyl 4-(2-pyridin-4- ylquinazolin-4-yl)piperazine-1- · carboxylate	+0-N-N-N-N-N-N-N-N-N-N-N-N-N-N-N-N-N-N-N

Table 3

#	Name	Structure
15	2-pyridin-4-yl-N-[(2S)-1,2,3,4- tetrahydronaphthalen-2-yl]quinazolin-4- amine	HN N N N N N N N N N N N N N N N N N N
16	4-[(1S)-2,3-dihydro-1H-inden-1- ylmethyl]-2-pyridin-4-ylquinazoline	
17	(1R,2S)-1-[(2-pyridin-4-ylquinazolin-4-yl)amino]-2,3-dihydro-1H-inden-2-ol	HO NO
18	(1S,2R)-1-[(2-pyridin-4-ylquinazolin-4-yl)amino]-2,3-dihydro-1H-inden-2-ol	HO,,,
19	1,1-dimethylethyl 4-[(2-pyridin-4- ylquinazolin-4-yl)amino]piperidine-1- carboxylate	HN N
20	2-pyridin-4-yl-N-{[2,4,6- tris(methyloxy)phenyl]methyl}quinazolin- 4-amine	N N N N N N N N N N N N N N N N N N N

Table 3

#	Name	Structure
21	N-piperidin-4-yl-2-pyridin-4-ylquinazolin- 4-amine	HIN—NH
22	N-{(1S,2S)-2- [(phenylmethyl)oxy]cyclopentyl}-2- pyridin-4-ylquinazolin-4-amine	N HN M
23	N-phenyl-N'-(2-pyridin-4-ylquinazolin-4-yl)benzene-1,4-diamine	
24	3-[(2-pyridin-4-ylquinazolin-4- yl)amino]naphthalen-2-ol	HO Z Z Z Z Z
25	N-{4-[(1-methylethyl)oxy]phenyl}-2- pyridin-4-ylquinazolin-4-amine	HH Z Z
26	(1S,2R)-1-[(2-phenylquinazolin-4-yl)amino]-2,3-dihydro-1H-inden-2-ol	HN—N N= N= N= N= N= N= N= N= N= N= N= N= N=

Table 3

#	Name	Structure
27	(1R,2S)-1-[(2-phenylquinazolin-4-yl)amino]-2,3-dihydro-1H-inden-2-ol	HN—N OH
28	(1R,2R)-2-[(2-phenylquinazolin-4- yl)amino]cyclopentanol	N N NH OH
29	(1R,2R)-2-[(2-phenylquinazolin-4- yl)amino]cyclohexanol	N NH OH
30	(1S,2R,3R,5R)-3-(hydroxymethyl)-5-[(2-phenylquinazolin-4-yl)amino]cyclopentane-1,2-diol	HO,,,, OH
31	(1S,2R)-1-[(6-chloro-2-pyridin-4- ylquinazolin-4-yl)amino]-2,3-dihydro-1H- inden-2-ol	HO
32	N-(2-piperazin-1-ylethyl)-2-pyridin-4- ylquinazolin-4-amine	N—NH N—NH

Table 3

#	Name	Structure
33	(1S,2R)-1-[(2-pyridin-3-ylquinazolin-4-yl)amino]-2,3-dihydro-1H-inden-2-ol	HN-N-N-N-N-N-N-N-N-N-N-N-N-N-N-N-N-N-N-
34	(1R,2S)-1-[(2-pyridin-3-ylquinazolin-4-yl)amino]-2,3-dihydro-1H-inden-2-ol	OH NOH NO
35	(1R,2R)-2-[(2-pyridin-3-ylquinazolin-4-yl)amino]cyclopentanol	N NH OH
36	(1R,2R)-2-[(2-pyridin-3-ylquinazolin-4- yl)amino]cyclohexanol	N N N OH
37	(1S,2R)-1-[(2-pyridin-2-ylquinazolin-4-yl)amino]-2,3-dihydro-1H-inden-2-ol	HN—N N—N N—N
38	(1R,2S)-1-[(2-pyridin-2-ylquinazolin-4-yl)amino]-2,3-dihydro-1H-inden-2-ol	HN—N—N

Table 3

#	Name	Structure
39	(2S)-3-[(2-pyridin-4-ylquinazolin-4-yl)amino]propane-1,2-diol	N OH OH
40	[(2S)-1-(2-pyridin-4-ylquinazolin-4-yl)- 2,3-dihydro-1H-indol-2-yl]methanol	HO III N
41	(2R)-2-[(2-pyridin-4-ylquinazolin-4- yl)amino]propan-1-ol	HO NH
42	(2S)-1-[(2-pyridin-4-ylquinazolin-4- yl)amino]propan-2-ol	N NH OH
43	(1S,2R)-1-{[2-(2-ethylpyridin-4-yl)quinazolin-4-yl]amino}-2,3-dihydro-1H-inden-2-ol	HO Z = Z
44	(1R,2S)-1-{[2-(2-ethylpyridin-4-yl)quinazolin-4-yl]amino}-2,3-dihydro-1H-inden-2-ol	HO Z Z

Table 3

#	Name	Structure
45	(1S,2R)-1-[(6-bromo-2-pyridin-4- ylquinazolin-4-yl)amino]-2,3-dihydro-1H- inden-2-ol	HO///NH N NH Br
46	(1S,2R)-1-{[6,7-bis(methyloxy)-2-pyridin-4-ylquinazolin-4-yl]amino}-2,3-dihydro-1H-inden-2-ol	HO:::\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\
47	1-(2-pyridin-4-ylquinazolin-4- yl)piperidin-3-ol	N—NOH
48	(1S,2R)-1-{[2-pyridin-4-yl-7- (trifluoromethyl)quinazolin-4-yl]amino}- 2,3-dihydro-1H-inden-2-ol	HO/// N NH
49	(1S,2R)-1-({2-[6-(methyloxy)pyridin-3-yl]quinazolin-4-yl}amino)-2,3-dihydro-1H-inden-2-ol	HN-N N-N-N-N-N-N-N-N-N-N-N-N-N-N-N-N-N-N

Table 3

#	Name	Structure
50	N-[(3S)-piperidin-3-yl]-2-pyridin-4- ylquinazolin-4-amine	N—NH NH
51	(1S,2R)-1-[(7-methyl-2-pyridin-4- ylquinazolin-4-yl)amino]-2,3-dihydro-1H- inden-2-ol	HN—N N—N
52	(1S,2R)-1-({2-[2,4-bis(methyloxy)pyrimidin-5-yl]quinazolin-4-yl}amino)-2,3-dihydro-1H-inden-2-ol	in OH OH NO
53	(2R)-3-methyl-2-[(2-pyridin-4-ylquinazolin-4-yl)amino]butan-1-ol	N OH NH
54	(2S)-3-methyl-2-[(2-pyridin-4- ylquinazolin-4-yl)amino]butan-1-ol	N—NH OH
55	(2S)-2-phenyl-2-[(2-pyridin-4- ylquinazolin-4-yl)amino]ethanol	N OH NH

Table 3

#	Name	Structure
56	(2R)-2-phenyl-2-[(2-pyridin-4- ylquinazolin-4-yl)amino]ethanol	N OH NH
57	(1S,2R)-1-[(2-pyridin-4-ylpyrimidin-4-yl)amino]-2,3-dihydro-1H-inden-2-ol	HN—N N—N N—N
58	(1S,2R)-1-[(2-pyrazin-2-ylquinazolin-4-yl)amino]-2,3-dihydro-1H-inden-2-ol	HN—N—N
59	(1S,2R)-1-{[2-(4-aminopyridin-3-yl)quinazolin-4-yl]amino}-2,3-dihydro-1H-inden-2-ol	HN-N N-NH ₂
60	(2R)-3-phenyl-2-[(2-pyridin-4- ylquinazolin-4-yl)amino]propan-1-ol	HO NATIONAL PROPERTY OF THE PR
61	(2S)-3-phenyl-2-[(2-pyridin-4- ylquinazolin-4-yl)amino]propan-1-ol	HO ZH Z

Table 3

#	Name	Structure
62	2-[(phenylmethyl)(2-pyridin-4- ylquinazolin-4-yl)amino]ethanol	N OH
63	(1S,2R)-1-{[2-(2-aminopyrimidin-4-yl)quinazolin-4-yl]amino}-2,3-dihydro-1H-inden-2-ol	HN-N N-N N-N H ₂ N
64	5-(4-{[(1S,2R)-2-hydroxy-2,3-dihydro- 1H-inden-1-yl]amino}quinazolin-2- yl)pyridin-2-ol	HN—N N N OH
65	(1S,2R)-1-({2-[2-(methylthio)pyrimidin-4-yl]quinazolin-4-yl}amino)-2,3-dihydro-1H-inden-2-ol	HN-N-N-N-N-N-N-N-N-N-N-N-N-N-N-N-N-N-N-
66	2-{4-[(2-pyridin-4-ylquinazolin-4-yl)amino]piperazin-1-yl}ethanol	OH ZZZZZZZZZZZZZZZZZZZZZZZZZZZZZZZZZZZZ

Table 3

#	Name	Structure
67	N-piperidin-1-yl-2-pyridin-4- ylquinazolin-4-amine	

- 19. A pharmaceutical composition comprising the compound according to any one of claims 1 18 and a pharmaceutically acceptable carrier.
- 20. A metabolite of the compound or the pharmaceutical composition according to any one of claims 1-18.
 - 21. Use of a compound according to any of claims 1 18 in the preparation of a medicament for modulating the *in vivo* activity of a kinase, for treating diseases or disorders associated with uncontrolled, abnormal, and/or unwanted cellular activities, or for inhibiting proliferative activity in a cell.
- 10 22. The use according to claim 21, wherein the kinase is Tie-2.
 - 23. The use according to claim 22, wherein modulating the *in vivo* activity of Tie-2 comprises inhibition of Tie-2.
- 24. A method of screening for a modulator of a Tie-2 kinase, the method comprising combining either a composition comprising at least one of the compound according to any of claims 1-18 and the pharmaceutical composition according to claim 31, and at least one candidate agent and determining the effect of the candidate agent on the activity of said kinase.

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